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# VERIFICATION OF TRANSLATION

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I verify that the attached English translation of International Patent Application Number PCT/FR02/02086 filed on 17th June 2002 is a true and correct translation made by me of the attached document in the French language;

I further declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that wilful false statements and the like so made are punishable by fine or imprisonment or both under Section 1001 of Title 18 of the United States code and that such wilful false statements may jeopardize the validity of the application or any patent issued thereon.

Date: 3rd December 2003 Lesley Stone

# GENE ASSOCIATED WITH LEISHMANIA PARASITE VIRULENCE

The invention relates to the field of the fight against leishmaniasis. It results from the identification, from wild isolates of Leishmania major, of a gene coding for a protein, designated LmPDI, having two regions identical to the sequence (Cys-Gly-His-Cys) of the potential active site of protein disulfide-isomerase (PDI). This LmPDI protein is predominantly expressed in the most virulent isolates of the parasite. It firstly constitutes a novel therapeutic target for developing anti-leishmaniasis drugs, and secondly, a novel element that can form part of the composition of immunogenic and possibly vaccinating preparations intended to protect a human or animal host against leishmaniasis.

Leishmaniases constitute a heterogeneic group of diseases that affect several million individuals and are due to infection of the host by a protozoic parasite of the genus Leishmania. Clinical expression of the infection is characterized by a high degree of polymorphism, including asymptomatic infection, simple or recurring cutaneous forms, diffuse or anergic cutaneous forms, mucocutaneous forms and visceral forms, which are fatal in the absence of specific treatment. In general, and depending on the geographical distribution of the disease, each species or sub-species of leishmaniasis is responsible for a particular clinical form; however, this is not a strict rule. Further, in the same geographical region, the same parasitic species can be responsible for clinical forms of varying severity. This diversity in clinical expression of the infection is at least partially due to a diversity in the virulence of the parasite.

During its cycle, the parasite alternates between two stages: the flagellate promoastigote stage, which is found in the digestive tract of the insect vector, and the amastigote stage in the host macrophage. Anti-leishmaniasis drugs are difficult to use, not least because they are toxic and because of the ever more frequent resistance developed by the parasite (Lira, Sundar et al, 1999). Further, recently developed and tested vaccines have so far not shown the expected efficacy (Sharifi, FeKri et al, 1998; Khalil, El Hassan et al, 2000).

The absence of tools for controlling leishmaniasis is partially explained by the complexity of the parasite transmission cycles and by the dearth of current knowledge regarding the biology of the parasite. During the last ten years, several molecules playing a fundamental role in the biology and infectivity of the parasite have been identified. Modifications to surface glyco-conjugates, particularly lipophospho-glycane (LPG), are associated with modifications to the infectivity and virulence of the parasite Leishmania (L) major and L. donovani (Beverley and Turco, 1998); Desjardins and Descoteaux, 1998; Sacks, Modi et al, 2000; Spath, Epstein et al, 2000), which does not appear to be the case for L. mexicana (Ilg 2000; Ilg, Demar et al, 2001). Molecules involved in the biosynthesis of LPG: phosphomannose isomerase (Garami and Ilg, 2001), LPG1 (Sacks, Modi et al, 2000; Spath, Epstein et al, 2000), LPG2 (Descoteaux, Luo et al, 1995) and galactosyl transferase (De and Roy, 1999) have also been associated with the virulence of Leishmania. Other factors in virulence have recently been described. They include the family of cysteine proteases (Mottram, Brooks et al, 1998), mitogen activated protein (MAP)-kinases (Wiese, 1998), the A2 gene (Zhang and Matlashewski, 1997), the surface glycoprotein gp63 (Chakrabarty, Mukherjee et al, 1996), kinetoplastid membrane protein (KMP)-11 (Mukhopadhyay, Sen et al, 1998), superoxide dismutase (Paramchuk, Ismail et al, 1997), trypanothione reductase (Dumas, Ouellette et al, 1997), and certain members of the heat shock protein (HSP) family (Hubel, Krobitsch et al, 1997).

Characterizing virulence factors may have fundamental implications in the development of novel drugs or vaccines against these diseases. Preferential screening of a protein involved in the virulence of a parasite can avoid the unnecessary appearance of resistance in less dangerous strains, which resistances may then be transmitted to other strains. Further, a mutation of the targeted virulence protein causing resistance to the drug can in that case also cause a reduction or even loss of the virulence of the parasite and thus have a certain therapeutic effect.

Over the past decades, a number of approaches have been used to study the virulence factors for the Leishmania parasite. These approaches were based on genetic studies, such as the

complementation of mutated parasites (Ryan, Garraway et al, 1993; Descoteaux, Luo et al, 1995; Desjardins and Descoteaux, 1997; Wiese, 1998), the use of the gene invalidation technique (Titus, Gueiros-Filho et al, 1995; Mottram, Souza et al, 1996; Dumas, Ouellette et al, 1997; Hubel, Krobitsch et al, 1997; Mottram, Brooks et al, 1998), or the analysis of genes for resistance to drugs on parasites manipulated in the laboratory (Cotrim, Garrity et al, 1999; Perez-Victoria, Perez-Victoria et al, 2001). Those studies identified several genes that were important to the biology of the parasite, and are currently being validated as targets for novel drugs (Selzer, Chen et al, 1997; McKerrow, Engel et al, 1999), or for the development and use of attenuated mutants as live vaccines (Titus, Gueiros-Filho et al, 1995; Streit, Recker et al, 2001). It is important to point out almost all of the studies carried out up to now on the virulence of the parasite Leishmania are based either on laboratory clones which have lost their virulence after prolonged culture, or on parasites genetically manipulated by mutagenesis experiments, gene invalidation or gene overexpression. Thus, it is possible that the conclusions relating to the virulence of the genes identified under these conditions are not actually relevant to the natural pathogenicity of the parasite in the transmission regions.

With the aim of studying the molecular bases for the virulence of the parasite, avoiding the methodological bias linked to the use of laboratory strains, the inventors initially isolated wild strains of Leishmania (L) major with different levels of virulence. L. major is the agent in zoonotic cutaneous leishmaniasis (ZCL), which exists in epidemic proportions in man over a very wide area which extends seamlessly from Mauritania to Mongolia. The inventors identified isolates of L major obtained from human ZCL lesions, all obtained during the same transmission season, and which differ in their pathogenicity in an experimental model of the infection in BALB/c sensitive mice (Example 1). The differences in experimental pathogenic power correlate with the differences in growth in vitro, which reflects the variations in the biology of these wild isolates.

The "differential display" technique (Liang and Pardee, 1992; Liang, Bauer et al, 1995) was then used to identify genes differentially expressed between completely different isolates through their experimental pathogenic power in the BALB/c mouse (two highly virulent isolates and two other less virulent isolates). This technique allows genes which are expressed at different levels to be studied without knowing their sequence in advance. Three transcripts that are preferentially expressed in the two most virulent isolates were then identified. One of these transcripts was completely characterized by dint of screening a cDNA library of L. major. An analysis of the sequence demonstrated a homology with the protein disulfide-isomerase family (PDI, Erp60 and Erp72) in eukaryotes. This novel protein has been designated LmPDI for the following reasons: like other members of the PDI family, (i) LmPDI possesses two CGHC active regions, (ii) the N-terminal region of this protein contains in a potential signal sequence and, in the carboxy-terminal region, a potential signal for retention in the endoplasmic reticulum (EEDL); (iii) it can organize itself into an oligomeric structure; (iv) the recombinant protein produced in E. coli expresses PDI activity in vitro. Further, outside the conserved regions mentioned above, there are very few similiarities between LmPDI and the other PDIs described above. In fact, the PDI family includes a plurality of highly divergent molecules involved in the maturation of proteins secreted into the endoplasmic reticulum (Noiva 1999; Frand, Cuozzo et al 2000). PDIs are multi-functional proteins which are involved in complex mechanisms of retention, repair, regulation of expression; they assist changes in conformation to allow only correctly folded proteins to leave the endoplasmic reticulum. In addition to their enzymatic functions (reduction and isomerisation), other functions have recently been attributed to PDIs; they include chaperone activities, the binding of peptides and cellular adhesion (Ferrari and Soling, 1999). It is important to emphasize that LmPDI is predominantly expressed in the most virulent isolates (Example 2). In total, these results suggest that LmPDI plays an important role in the natural virulence of the Leishmania parasite, and can thus constitute a novel target for chemotherapy or vaccination.

Further, recent data regarding the involvement of the bacterial equivalent of PDIs (Martin, 1995; Ostermeier, De Sutter et al, 1996) (termed DsbA, disulphide bond) make suggestions regarding the role this protein could play in the pathogenicity of different micro-organisms (Yu and Kroll, 1999). Deactivation of the DsbA gene dramatically affects the survival and virulence of Shigella flexneri (Yu, 1998; Yu, Edwards-Jones et al, 2000). DsbA is also involved in the genesis of the enterotoxin of Vibrio cholerae (Peek and Taylor, 1992; Yu, Webb et al, 1992). DsbA also important for the pathogenicity of pathogenic Escherichia coli species: in species that are pathogenic for the urinary tract, it catalyses the formation of disulfide bridges of a specific chaperone protein of pilin (Zhang and Donnenberg, 1996). In enteropathogenic species, it is also required for the stability and formation of pili (Hultgren, Abraham et al, 1993; Wang, Bjes et al, 2000).

The present application constitutes the first description suggesting an important role for PDIs as a virulence factor in a protozoic parasite. LmPDI could exert its effects by assisting changes in the conformation and stability of other factors essentially to the biology and pathogenicity of the parasite Leishmania. Identifying such factors would be extremely advantageous for a better comprehension of the biology of this parasite, and for the development of novel treatments or vaccines against leishmaniases.

Thus, in a first aspect the invention concerns a protein involved in the virulence of Leishmania, comprising at least one (Cys-Gly-His-Cys) site identical to the potential active site of a protein from the protein disulfide-isomerase family (PDI). This protein is preferably a protein coded by the parasite itself.

In particular, the invention concerns the LmPDI protein of Leishmania major, with sequence SEQ ID No: 3, and any functional variant of LmPDI having at least 40% identity, preferably at least 80% identity with LmPDI.

We shall define here a "functional variant of LmPDI" as a protein that is capable of complementing LmPDI in an infectivity test carried out with a strain of L. major in which the

LmPDI gene has been deactivated. The term "infectivity test" as used here means any test that can evaluate the biological properties relating to the growth of the parasite conventionally associated with virulence. In particular, the three following types of tests can be cited:

- growth kinetics in a liquid medium of the promastigote form of the parasite, for example by a technique of the type described in point 2 of Example 5;
- the infection capacity of mouse macrophages cultivated in vitro, using a technique such as that described in Example 6 and in the article by Kebaïer et al, 2001;
- the capacity to induce experimental murine leishmaniasis by infecting sensitive mice (BALB/c, for example). The technique is detailed in point 3 of Example 5, for example, and in the article by Kebaïer et al, 2001.

The percentage of identity with LmPDI are evaluated using CLUSTAL W version 1.8 software (Thompson J D, Higgins D G and Gibson T J) or BOXSHADE version 3.21 software (Hoffman K and Baron M) which produced percentage identities of LmPDI with proteins from the PDI family of several species as between 27% and 36% (Example 2).

In a second aspect, the invention concerns a recombinant polypeptide comprising at least one fragment of more than 10 amino acids of a protein as defined above, if appropriate fused with a further polypeptide fragment, said recombinant polypeptide being capable of triggering an immunological reaction against an epitope of LmPDI when administered to an animal. The invention also concerns a recombinant polypeptide comprising at least one fragment of more than 10 amino acids of a protein as defined above, if appropriate fused with a further polypeptide fragment, said recombinant polypeptide being capable of being recognized by antibodies directed against the LmPDI protein.

Throughout this text, the term "polypeptide" should be taken in its broad sense, i.e., including sequences of at least 10 amino acids (or more when stated), which may or may not comprise glycosylated motifs or glycolipids, and regardless of its primary, secondary or tertiary

structure. The LmPDI fragment present in the recombinant polypeptides of the invention described above may be over 15, 20, 30, 50 or 100 amino acids in size, or even more.

The LmPDI, recombinant or purified from infected cells, and a polypeptide of the invention can be used to immunize a human or animal host, to protect it from leishmaniasis or to produce and recover antibodies directed against LmPDI, as described in Example 2.

A particular recombinant polypeptide of the invention is the LmPDI-$(His)_6$ protein with sequence SEQ ID No: 4, described in Example 2.

A further example of the recombinant polypeptide of the invention is a fusion protein between a LmPDI fragment comprising at least one epitope of LmPDI and a carrier polypeptide contributing to the presentation of that fragment to the immune system. It can in particular be a fusion of all or a portion of the LmPDI with a fragment of β-lactamase, or a tetanus or diphtheria anatoxin, or any other polypeptide from a pathogenic organism, in particular of parasitic, bacterial or viral origin.

In a further aspect, the invention concerns a nucleic acid sequence coding for a protein or a polypeptide as described above. A preferred nucleic acid sequence comprises the sequence coding for LmPDI with sequence SEQ ID No: 2, or a fragment of said sequence with a size of 30 nucleotides or more, preferably more than 100 nucleotides, coding for a polypeptide comprising at least one characteristic epitope of LmPDI.

The invention also concerns a nucleic acid vector comprising a nucleic acid sequence of the invention. As an example, it may be a plasmid, a cosmid, a phage or a virus. Preferably, a vector of the invention will allow expression in a host cell of a protein or a polypeptide in accordance with the invention. In particular, a vector of the invention can allow expression of LmPDI in a bacterial or eukaryotic cell.

The invention also pertains to a cultured cell comprising a vector as defined above. Said cell can be a bacterium, a yeast, an insect cell, a mammalian cell or any other type of cell. It can be used either to express and possibly produce a protein or a polypeptide in accordance with the

invention, or to produce a vector which will then serve to express a protein or a polypeptide in accordance with the invention in a further cultured cell type, or in vivo. Purely by way of non-limiting illustration, CHO, VERO, BHK21 cells and insect cells can be cited as cell types that can be used in vitro in the context of the present invention. Similarly, BCG and Salmonella typhimurium can be cited as cells that can be used in vivo. Finally, it is important to note that administration to an individual of a viral vector, for example a vaccine virus or DNA coding for a polypeptide or a protein as described above for vaccine purposes is also encompassed within the scope of the invention.

A particular cell of tne invention is the bacterial strain LmPDI-$XL_1$ deposited at the Collection National de Culture des Microorganismes [CNCM, the National Collection of Microorganism Cultures], on 31/01/2002 with accession number I-2621. This strain is derived from a XL1-blue MRF' strain bacterium with genotype Δ(mrcA)183 Δ(mcrCB-hsdSMR-mrr)173endA1 sup E44 thi-1 recA1 gyrA96 relA1 lac[F' proAB $lac^qZ$ ΔM15 Tn10 (Tet')], transformed by the plasmid pBK-CMV-LmPDI. This plasmid corresponds to the plasmid pBK-CMV sold by Stratagene (La Jolla, CA) to which cDNA from LmPDI has been added between the EcoRI and Xho I restriction sites.

The invention also pertains to a nucleic acid probe which specifically hybridizes under stringent conditions with the nucleic acid sequence of SEQ ID No: 2, allowing the presence or absence of the virulence gene coding for LmPDI to be determined in a biological sample.

"Stringent hybridization conditions" are defined herein as conditions that allow specific hybridization of two DNA molecules at about 65°C, for example in a solution of 6X SSC, 0.5% SDS, 5X Denhardt's solution and 100 µg/ml of denatured non specific DNA or any solution with an equivalent ionic strength, and after a washing step carried out at 65°C, for example in a solution of at most 0.2X SSC and 0.1% SDS or any solution with an equivalent ionic strength. However, the stringency of the conditions can be adapted by the skilled person as a function of the size of the sequence to be hybridized, its GC nucleotide content, and any other parameter, for

example following protocols described by Sambrook et al, 2001 (Molecular Cloning: A Laboratory Manual, 3rd Edition, Laboratory Press, Cold Spring Harbor, New York).

In the above definition, and throughout the present text, the term "specific" should be taken to have its broadest meaning, normally used in laboratories. Thus, a molecule A specifically recognizes a molecule B if, in a complex mixture, molecule A has an affinity for molecule B that is significantly higher than its affinity for other molecules of the mixture, so that it is possible to detect molecule B via molecule A.

The stringency conditions used here are those that allow the PDIs of different Leishmania species to be detected rather than those of the host and other microorganisms in the presence of a radiolabelled probe synthesized from the cDNA of LmPDI.

As an example, a probe of the invention, which specifically hybridizes with sequence SEQ ID No: 2 under stringent conditions, is such that a Southern blot carried out using said labeled probe, when carried out on a DNA sample from cells infected with a strain of L. major expressing LmPDI, has at least one clearly distinct band of higher intensity than other bands (non specific), said band not appearing on a Southern blot carried out under the same conditions on a DNA sample from cells not infected by a strain of L. major.

In a further aspect, the invention concerns a nucleotide primer that can allow specific amplification of at least a portion of the sequence SEQ ID No: 1, from cells infected with Leishmania, thus allowing the presence or absence of the virulence gene coding the LmPDI to be determined in a biological sample. Amplification will be termed "specific" if the amplification reaction carried out from control cells not infected with Leishmania does not result in significant amplification of any sequence, while the same reaction carried out on a sample containing the nucleotide sequence of SEQ ID No: 1 results in amplification of at least one fragment of said sequence.

The probes and primers mentioned above can if necessary be labeled and/or presented in diagnostic kits which also form part of the invention. It may be advantageous to determine the

presence and possibly the level of expression of the gene for LmPDI during an infection with Leishmania, for example to determine the parasitic and/or opportunistic charge of a treatment involving the use of a LmPDI inhibitor.

In a further implementation, the invention provides purified antibodies specifically recognizing LmPDI. They may be monoclonal or polyclonal human, humanized or animal antibodies. Said antibodies can be purified, for example, on an LmPDI affinity column using the protocol described in the experimental section. Said specific LmPDI antibodies may have a number of applications.

They may serve to detect the presence of LmPDI in a biological sample, for example to diagnose leishmaniasis and/or to determine the possibility of using a LmPDI inhibitor to treat that leishmaniasis.

Thus, the invention also concerns an in vitro method for diagnosing an infection by a parasite responsible for leishmaniasis. Such a method can be carried out using a polypeptide or a protein of the invention or an antibody directed against that protein, or using probes as defined above.

A particular diagnostic method of the invention comprises the following steps:

- bringing at least one antibody in accordance with the invention into contact with a biological sample from a subject partially infected by a parasite responsible for leishmaniasis under conditions allowing the formation of an immune complex between said antibody and antigenic proteins contained in the sample;
- detecting said complex.

The complex can be detected using any means that is known to the skilled person (enzymatic reaction, fluorescence transfer or the like).

The antibodies of the invention can be comprised in diagnostic kits in the same manner as the probes or primers mentioned above.

Diagnostic kits for implementing the method described above form an integral part of the present invention.

By way of example, such a kit can comprise

- at least one antibody in accordance with the invention;
- a medium suitable for forming an immune complex between the antigenic proteins contained in the analyzed sample and said antibody;
- reagents allowing the detection of the complexes so formed;
- if appropriate, control samples.

Alternatively, the antibodies of the invention can form part of the composition of a drug intended for prophylaxis, attenuation or for the treatment of certain leishmaniases.

In a further aspect of the invention, the invention pertains to an immunogenic composition comprising a protein and/or a recombinant polypeptide and/or a nucleic acid sequence and/or a vector and/or a cell of the invention as described above, said immunogenic composition being capable of in vitro stimulation of the proliferation of mononuclear cells deriving from individuals who have come into contact with a Leishmania parasite. A preferred immunogenic composition of the invention is capable of in vitro stimulation of the proliferation of mononuclear cells deriving from individuals who have come into contact with Leishmania major.

In a preferred implementation of the immunogenic compositions of the invention, said compositions have a formulation that is pharmaceutically acceptable for administration to a human or animal host.

The inventors have shown that LmPDI is susceptible of in vitro induction of the production of cytokines by mononuclear cells deriving from individuals who have come into contact with L. major, and that the expression profile of the cytokines corresponds to that observed during a type Th1 immune response (Example 3). An immunogenic composition as described above, which is capable of inducing a type Th1 immune response when administered

to a human or animal host, thus constitutes a particularly preferred implementation of the present invention.

The invention also pertains to a vaccine composition comprising a protein and/or a recombinant polypeptide and/or a nucleic acid sequence and/or a vector and/or a cell of the invention as described above, said vaccine composition being intended to protect a human or animal host against leishmaniasis. Preferably, the vaccine compositions of the invention are formulated in a manner that is pharmaceutically acceptable for administration to a human or animal host.

Said vaccine composition can be in the liquid form for injection into a patient, either subcutaneously or intramuscularly, or in the form of an oral vaccine, in the form of a pomade, or in the form of particles bound to a nucleotide sequence of the invention, for example by DNA adsorption onto the particle surface. This latter form allows the vaccine to be administered using a gene gun. It is important to note that the formulations for the vaccine compositions mentioned here are given solely by way of example and are in no way restrictive.

The immunogenic and/or vaccine compositions of the invention can also comprise one or more antigen(s) that are heterologous as regards Leishmania, and/or one or more nucleic acid sequence(s) coding for said antigens. The compositions of the invention can thus trigger an immunological reaction against several different pathogens and if appropriate may constitute polyvaccines.

The vaccination process and the doses of active agent must be adapted to the type of vaccine used and to the mammal to which it is administered.

Methods for vaccination against Leishmania, consisting of administering a composition comprising a protein and/or a recombinant polypeptide and/or a nucleic acid sequence and/or a vector and/or a cell of the invention as described above to a human or animal host, are also encompassed by the invention.

Determining the role of LmPDI in the virulence of Leishmania can also enable novel strategies for identifying active molecules for inhibiting the growth of the parasite to be envisaged. It has been shown that a molecule inhibiting PDI, for example, such as bacitracin or chloromercuribenzene sulfonic acid (pCMBS), inhibits the growth of Leishmania in a liquid medium (Example 4). Thus, the invention also pertains to a method for screening molecules susceptible of inhibiting the growth of Leishmania major, comprising a step for evaluating the capacity of said molecules to inhibit the activity of LmPDI. Protein disulfide-isomerases in general have a plurality of activities, in particular oxido-reduction, isomerase, and chaperone activities. The screening methods of the invention can pertain to the inhibition of any of the functions of LmPDI.

In a particular screening method of the invention, the step for evaluating the capacity of a molecule to inhibit the activity of LmPDI is carried out in a test for reactivating scrambled RNase A, comprising the following steps:

- incubating scrambled RNase A in the presence of LmPDI under conditions allowing its reactivation;
- incubating scrambled RNase A under conditions identical to those allowing its reactivation by LmPDI, the molecule to be tested being added;
- comparing the results obtained in the absence and in the presence of the test molecule, a fault in the reactivation of RNase A in the presence of the test molecule revealing that said molecule has an LmPDI inhibiting activity.

Any other PDI activity test can be used in the screening methods of the invention, in particular any test derived from the initial protocol described by Lyles and Gilbert (1991).

A screening method of the invention can also comprise a test for inhibiting the growth of Leishmania major in a liquid medium and if appropriate, a test for inhibiting the growth of Leishmania major in an experimental murine model of leishmaniasis. An example of such a method is described in the experimental section, Example 5.

The active molecules screened by the method defined above are characterized by their capacity to inhibit or modulate the growth of Leishmania major.

The results obtained with bacitracin shown in Example 4 show that a PDI inhibitor can inhibit the growth of Leishmania. The use of one or more protein disulfide-isomerase (PDI) inhibitors for the preparation of a pharmaceutical composition intended for prophylaxis, attenuation, or treatment of an infection with Leishmania thus forms an integral part of the invention. Compounds with an anti-PDI activity that can be used in accordance with the invention that can be cited are anti-PDI or anti-LmPDI antibodies, bacitracin, zinc bacitracin, 5,5'-dithiobis(2-nitrobenzoic) acid (DTNB), p-chloromercuribenzene sulfonic acid (pCMBS) or tocinoic acid.

The compositions prepared in accordance with the above uses can preferably be topically, orally or parenterally administered to a human or animal host.

In accordance with a particular aspect, the invention concerns the use of bacitracin or zinc bacitracin as an inhibitor to the growth of a parasite responsible for leishmaniasis or as an active agent against a Leishmania infection.

Clearly, a pharmaceutical composition for the treatment of an infection with Leishmania containing one or more protein disulfide-isomerase (PDI) inhibitors forms an integral part of the invention. Such a composition can in particular contain bacitracin or zinc bacitracin. The composition can be formulated for topical application, for example in the form of a cream, an ointment, a pomade, or a spray, this list being non-limiting. The inventors have shown that such a composition, applied locally in the form of a pomade to the injection site of the parasite in BALB/c mice, attenuates the progress of the disease (Example 9 and Figure 14).

The present invention also pertains to a pharmaceutical composition for the treatment of an infection with Leishmania, comprising at least one specific antibody for LmPDI and/or any molecule that inhibits PDI activity. Such a composition is preferably appropriate for topical, oral or parenteral administration.

Methods for treating leishmaniases, comprising administration of a PDI or LmPDI inhibitor to a human or animal patient, whether an antibody or any other type of molecule, also fall within the scope of the invention.

The examples and figures below describe the biological experiments which have been carried out in the context of the present invention and which provide the required experimental support, without in any way limiting its scope. They also illustrate, in a non restrictive manner, certain aspects of the implementation and importance of the present invention.

**KEY TO FIGURES**

Figure 1 shows a differential display (DD) analysis of the expression of Leishmania major genes in the two most virulent isolates (94 and 67, V) and the two least virulent isolates (32 and 07, v).

Figure 1A shows a portion of a sequencing gel after autoradiography, showing the products amplified by PCR using an arbitrary decamer and an oligo dT primer. The differentially expressed cDNAs are indicated by arrows. The p14 cDNA is indicated by an asterisk.

Figure 1B shows a Northern blot analysis of the expression of a gene identified by the DD technique between the most virulent isolates (94 and 67, V) and the least virulent isolates (32 and 07, v). The mRNA extracted from the promastigotes from different isolates in the stationary growth phase were hybridized with the radiolabelled probe p14. After autoradiography, the blots were de-hybridized then re-hybridized with a specific probe for the gene for the α-tubuline of L. Major (α-tub).

Figure 2 shows the nucleotide sequence for the cDNA (SEQ ID No: 1) of LmPDI and the deduced sequence of amino acids (SEQ ID No: 3). The nucleotides in lower case letters represent non-translated regions. The leader sequence (SL) of 18 nt is underlined and the potential sequence for the polyadenylation signal is boxed. The potential sequence for the

peptide signal is shown in bold. The potential active sites for LmPDI are double underlined and the probable sequence for retention in the endoplasmic reticulum is shown as a broken line.

Figure 3 shows the alignment of the amino acid sequence for LmPDI with the protein disulfide-isomerase of Trypanosoma brucei (T brucei, GenBank accession no.: P12865), Hypocrea jecorina (H. Jecorina, 074568), Caemorhabditis elegans (C. elegans), 017908), Chlamydomonas reinhardtii (C. reinhard, 048949), Drosophila melanogaster (D. melano, P54399), Cryptosporidium parvum (C. parvum, Q27553), and Homo sapiens (H, sapiens, P072237). The letters boxed in black indicate identical amino acids and those boxed in gray indicate similar amino acids. The "gaps" were introduced to obtain the maximum similarity between the aligned sequences and are indicated by dashes.

Two software programs were used to carry out the alignments:

- CLUSTAL W version 1.8; Thompson, J D, Higgins, D G and Gibson, T J;
- BOXSHADE version 3.21; Hoffman, K and Baron, M.

Figure 4 shows an analysis of the role of recombinant LmPDI in reactivating scrambled RNase. Scrambled RNase A (8 μM) was incubated in a buffer containing 4.5 mM (cCMP), 1 mM glutathione GSH, 0.2 mM glutathione disulfide GSSH, 2 mM EDTA and 100 mM Tris-HCl pH 8 in the presence of bovine serum albumin (BSA) (1.4 μM) as a negative control, bovine protein disulfide-isomerase (1.4 μM) as the positive control, or recombinant LmPDI (1.4 μM) for 30 minutes at 25°C. RNase A reactivation was determined by measuring the RNase A activity at 296 nm every 5 minutes for 30 minutes (Lyles and Gilbert, 1991).

Figure 5A shows a Southern Blot analysis for the number of copies of the LmPDI gene in the Leishmania major gene. 8 μg of isolate genomic 94 DNA from L. major was digested by the following restriction enzymes: AvaI, EcoRV, HindIII, PstI, EcoRI, XhoI, NcoI, SacI, SphI. The enzymes marked with an asterisk cleave once within the cDNA of LmPDI.

Figure 5B shows a Southern Blot analysis of the LmPDI gene in different species of Leishmania. 8 μg of genomic L. major DNA (94), dermotropic L. infantum (L. infantum MC),

viscerotropic L. infantum (L. infantum Visc); L. donovani were digested with the PstI enzyme. Genomic DNA was hybridized in these experiments by the probe representing the entire cDNA sequence of LmPDI.

Figure 6 shows immunodetection of native LmPDI in L. major with different preparations of anti-LmPDI antibodies. 20 µg of total promastigote GLC 94 proteins in Laemmli buffer (track 1) or in the presence of 0.5 m of DTT (track 2) and 0.05 µg of LmPDI produced in E. coli bacteria and purified (rLmPDI) (track 3) underwent electrophoresis then were transferred onto a nitrocellulose membrane, then revealed with an anti-LmPDI immunoserum (track 1) or anti-LmPDI antibodies purified on an affinity column (tracks 2 and 3).

Figure 7 shows a Western blot analysis of the expression of LmPDI in the two most virulent isolates (94, 67, V) and the two least virulent isolates (32, 7, v) from promastigotes. 20 µg of total promastigote protein in the stationary growth phase from different isolates underwent electrophoresis and were then transferred onto a nitrocellulose membrane which was incubated in the presence of polyclonal anti-LmPDI antibody. The arrows (>) indicate the 3 proteins recognized by the anti-LmPDI immunoserum.

Figure 8 shows the proliferation of mononuclear cells from individuals living in a zoonotic cutaneous leishmaniasis region in Tunisia, after incubation of LmPDI (5 µg/ml). The lymphomonocytary cells were recovered, washed by 3 successive centrifuge runs with RPMI-PS/Glu medium (30 ml then twice 10 ml) then counted and incubated at a concentration of $10^6$ cells/ml of medium in the presence or absence of a concentration of 5 µg/ml of LmPDI. After 5 days of culture, lymphocyte stimulation was estimated by incorporating tritiated thymidine. The result is expressed in CPM.

Figure 9 shows the results of a parasite (L. major) growth inhibition test in a liquid medium using bacitracin. They are growth curves taken over 96 hours, for promastigotes of L major in the presence of 0, 1 mM, 1.5 mM or 2 mM of bacitracin.

Figure 10 shows the effect of bacitracin (BAC), zinc bacitracin (BACZn), p-chloromercuribenzoic acid (pCMBA) and tocinoic acid (TOC) on the in vitro activity of recombinant LmPDI. Different concentrations of inhibitors (0 to 2 mM) were used to follow the effect of PDI inhibitors on the capacity of LmPDI to reactivate scrambled RNase A in vitro. LmPDI without inhibitors was used as the positive control (T).

Figure 11 shows the effect of bacitracin (BAC) (Figure 11A), zinc bacitracin (BACZn) (Figure 11B), 5,5'dithiobis-(2-nitrobenzoic acid) (DTNMB) (Figure 11C) and p-chloromercuribenzoic acid (pCMBA) (Figure 11D) on the in vitro growth of Leishmanias in a liquid medium. Different concentrations of inhibitors (0 to 5 mM) were used to follow the effect of PDI inhibitors on the multiplication of parasites in vitro. Parasites that had not been treated with inhibitors (T) were selected as a control for these experiments.

Figure 12 shows the inhibition of the activity of rLmPDI by bacitracin and zinc bacitracin. The effect of bacitracin (BAC) and zinc bacitracin (BACZn) on the activity of rLmPDI was measured in vitro. Different concentrations of BAC and BACZn inhibitors (0 to 2 mM) were tested to analyze their effect on the capacity of rLmPDI to reactivate scrambled RNase in vitro. The activity of rLmPDI in the absence of inhibitors acted as a positive control.

Figure 13 illustrates the inhibition of the multiplication of GLC94 promastigotes by zinc bacitracin. The effect of zinc bacitracin (BACZn) on the multiplication of GLC94 promastogotes was determined in vitro. Different concentrations of inhibitor were tested to analyze their effect on the capacity of the parasites to multiply in vitro. The control (C) was constituted by parasites cultivated in a complete medium in the absence of inhibitors.

Figure 14 shows the effect of zinc bacitracin on the evolution of the disease in sensitive BALB/c mice infected with GLC94 isolate promastigotes. Sensitive BALB/c mice were infected with $10^6$ promastigotes from the GLC94 isolate into the plantar pad and treated or not treated with bacitracin. The treatment was halted 9 weeks after infection (the arrow indicates treatment stoppage). Each curve shows the change in the size of a single mouse.

## EXAMPLES

The experimental results shown in the following examples were obtained using the following materials and methods:

Parasites and culture conditions

The L. major isolates used in this study derived from human ZCL lesions obtained during the study summarized in Example 1. The parasites were cultivated in NNN medium (solid medium prepared and based on agarose and rabbit blood) at 26°C, and progressively transferred into RPMI (SIGMA, St Louis, MO) containing 2 mM of L-glutamine, 100 U/ml of penicillin, 100 µg/ml of streptomycin and 10% of deactivated fetal calf serum (complete medium). Promastigotes in the logarithmic growth phase were adjusted to $10^6$ parasites/ml in a constant volume of complete medium and incubated at 26°C. The stationary growth phase was reached after 4 to 6 days with the density of parasites of $3 \times 10^7$ to $8 \times 10^7$ parasites. Those promastigotes in the stationary growth phase were used for RNA and protein extractions.

RNA extraction and Differential Display

Total RNA was extracted using the "TRIZOL" reagent (Gibco-BRL). PolyA$^+$ RNA was purified by passage through an oligo dT/cellulose column using a "poly A$^+$ RNA isolation kit" (Amersham-Pharmacia) following the manufacturer's instructions. 200 ng of mRNA was used in a reverse transcription reaction of 20 µl containing 1 µM of an oligo$(dT)_{11}$MN primer, with M = A or C or G and N = A or C or G or T (Genset), 1X First Strand Buffer (Gibco-BRL), 5 µM dNTP (Amersham-Pharmacia), 10 U of RNAsin (Promega) and 200U of reverse transcriptase (Gibco-BRL).

After incubating at 37°C for one hour, the reaction was stopped by incubating for 5 minutes at 95°C. The cDNA was amplified by PCR using a combination of 12 oligo dT and 10 arbitrary decamers. PCR was carried out in a volume of 20 µl containing 2 µl of reverse transcription reaction, 0.2 µM of 5' primer, 1 µM of 3' primer, 2 µM of dNTP, 10 µCi [$\alpha^{35}$S] of ATP, 1X Taq DNA polymerase reaction buffer and 1U of Taq polymerase (Amersham-

Pharmacia). The reactions were incubated in a Perkin-Elmer 9600 thermocycler for 40 cycles at 94°C for 30s, 40°C for 60s and 72°C for 30s followed by one cycle at 72°C for 6 minutes. The PCR products were analyzed on a 6% acrylamide sequencing gel. The gel was vacuum dried on Whatmann 3MM paper and autoradiographed. The differentially expressed cDNA was excised from the gel, eluted and reamplified by PCR in the presence of the same oligonucleotides, under the conditions described above. The amplification products were cloned in pMOSblue vector using the Blunt-ended PCR cloning kit (Amersham Pharmacia), following the manufacturer's instructions. The cloned fragments were sequenced using a Sequencing Ready Reaction Kit (Perkin-Elmer) and analyzed using the ABI 377 automatic sequencer.

Northern Blot analysis

200 ng of mRNA from promastigotes extracted during the stationary growth phase of 4 isolates from L major were denatured, separated on a 1.2% agarose /2.2 M formaldehyde gel and transferred by capillarity on a "Hybond $N^+$" (Amersham-Pharmacia) membrane. The nucleic acids were then fixed by heating for 2 hours at 80°C. The differentially expressed cDNA fragments and α-tubulin were labelled with [$\alpha^{32}$P]dCTP using the Megaprime DNA labelling system kit (Amersham-Pharmacia). Hybridizations were carried out in a 1X Denhardt's/6X SSC/0.1% SDS/0.1 mg.$ml^{-1}$ salmon sperm solution overnight at 65°C. The membranes were washed at 65°C in a solution containing 0.1X SSC/0.1% SDS and autoradiographed.

Construction of a cDNA library and characterization of LmPDI cDNA

A cDNA library was constructed from 5 µg of mRNA from promastigotes from the most virulent strain (GLC94) in the ZAPII vector, following the manufacturer's instructions (Stratagene). 6 x $10^6$ lysis plaques were screened using the p14 probe labeled with [$\alpha^{32}$P] dCTP using the Megaprime DNA labelling system kit (Amersham-Pharmacia). The lysis plaques of interest were removed and screened again to isolate positive clones from contaminating clones. The positive clones were then sequenced.

Southern Blot analysis

10 µg of genomic DNA extracted from promastigotes from the most virulent strain GLC94 were digested with the restriction enzymes indicated in Figure 5 and analyzed on a 0.6% agarose gel, then transferred to a Hybond $N^+$ (Amersham-Pharmacia) membrane. The membrane was incubated in the presence of a probe radioactively labeled with [$\alpha^{32}$P]dCTP and corresponding to the entire cDNA clone of LmPDI. The membranes were then washed in a solution containing 0.1X SSC/0.1% SDS and autoradiographed.

Expression and purification of the recombinant protein LmPDI in E. coli BL21 bacteria

The sequence corresponding to the open reading frame of cDNA of LmPDI (1371bp) deprived of the sequence coding for the peptide signal was cloned in the bacterial expression vector pET-22b (Novagen). E. coli BL21 bacteria containing the recombinant plasmid (pET-22b-LmPDI) were cultivated in LB medium then synthesis of the recombinant protein was induced in the presence of 1 mM of isopropyl-1-thio-D-galactopyranoside (IPTG) for 4 hours. The recombinant protein LmPDI-$(His)_6$ (SEQ ID No: 4) was purified by affinity chromatography on a nickel column ($Ni^{2+}$) (Amersham-Pharmacia). The purity of the protein produced was verified by SDS-PAGE.

Production of a polyclonal anti-LmPDI antibody and analysis of expression of the native protein by immunoblot

A rabbit was immunized by intramuscular injection of 500 µg of emulsified purified recombinant LmPDI in incomplete Freund's adjuvant (IFA, Sigma) (v/v). The rabbit received two additional injections of 500 µg of recombinant protein, the first intramuscularly 15 days after the first injection and the second intradermally 30 days later. The rabbit was bled 10 days after the last injection; the serum was harvested and kept at -80°C. The protein lysate from the promastigotes was denatured in Laemmli 1X buffer for 10 minutes at 100°C, deposited on a 12% SDS-acrylamide gel and electrotransferred onto a nitrocellulose membrane (Millipore). The membranes were incubated in a saturated PBS/0.1%Tween20/3% skimmed milk solution at ambient temperature for one hour, then in the same solution containing anti-LmPDI antibody

diluted to 1/1000$^{th}$ at 4°C overnight. After 3 washes in PBS/0.1%Tween20, the membranes were incubated in the presence of secondary rabbit anti-IgG antibody coupled with peroxidase (Amersham-Pharmacia, diluted to 1/1000) for one hour at ambient temperature and washed 3 times in PBS/0.1% Tween20. The protein-antibody complexes were revealed by detecting the peroxidase activity using the "ECL system" kit, following the manufacturer's instructions (Amersham-Pharmacia).

Preparation of scrambled RNase A

20 mg of purified ribonuclease (RNase A) was scrambled at ambient temperature for 18 hours in a buffer containing 0.15 M of DTT, 6 M of guanidine-HCl and 0.1 M of Tris-HCl at a pH of 8.6 before being purified on a Sephadex G-25 column equilibrated in 0.01 M HCl. The concentration of scrambled RNase A fractions was determined using an extinction coefficient of 9200 $M^{-1}cm^{-1}$ at 275 nm. The fractions were stored at -80°C for two weeks.

Reactivating RNase A in the presence of recombinant LmPDI protein

Scrambled RNase A (8 µM) was incubated in a buffer containing 4.5 mM cCMP, 1 mM glutathione GSH, 0.2 mM glutathione disulfide GSSH, 2 mM EDTA and 100 mM of Tris-HCl pH 8 in the presence of bovine serum albumin (BSA) (1.4 µM) as a negative control, bovine protein disulfide-isomerase (1.4 µM) as a positive control, or recombinant LmPDI (1.4 µM) for 30 minutes at 25°C. RNase A reactivation was determined by measuring the RNase A activity at 296 nm every 5 minutes, as described in the literature (Lyles and Gilbert, 1991).

**Example 1: Selection of L. major isolates having different levels of virulence**

The L. major isolates used in this study derived from human ZCL lesions obtained during a prospective study carried out in 1994-1995 at El Guettar, in southern Tunisia (Louzir, Melby et al, 1998). They had been selected from 19 isolates on the basis of their pathogenic power during experimental infection of sensitive BALB/c mice: 2 x $10^6$ amastigotes from various isolates were injected into the rear paw pads of BALB/c mice and the progress of the lesion was observed every week for 9 weeks. Five weeks after infection, the production of IL-4 and IFN-γ by

mononuclear cells of lymphatic ganglia activated in vitro by antigens from the parasite was measured.

These experiments showed firstly the great heterogeneity in the progress of the disease induced by different isolates of L. major and secondly, that using a single isolate leads to reproducible results.

The most virulent strains induced the greatest amount of IL-4 and the lowest levels of IFN-γ in vitro, 5 weeks after infection.

From the observation that clinical expression of infection with L. major varies depending on the strains and is reproducible within each of them in the experimental model of infection of BALB/c sensitive mice, the inventors devised the hypothesis that the genes involved in virulence could be differentially expressed when comparing the most virulent isolates with the least virulent isolates. A preliminary analysis of the expression of a group of genes already described by other authors and associated with the virulence of the parasite, including LPG1, LPG2, KMP-11, Cpc, Cpb, Hsp100, Gene B and gp63, was carried out by a reverse transcription technique and quantitative gene amplification. This analysis did not show any difference between L. major isolates expressing a different pathogenicity in BALB/c mice (Kebaier, Louzir et al, 2001).

Two isolates, MHOM/TN/94/GLC94 and MHOM/TN/94/GLC67 (GLC94 and GLC67 respectively), which induced severe lesions, developed rapidly and represented the most virulent isolates, and 2 isolates MHOM/TN/94/GLC07 and MHOM/TN/94/GLC32 (GLC07 and GLC32 respectively) inducing a less severe experimental disease and representing less virulent isolates, were selected to continue the search for virulence genes potentially expressed at different levels depending on the strains.

**Example 2: Differential Display identification of a novel protein disulfide isomerase LmPDI of Leishmania major, involved in natural parasite virulence**

1-Identification of genes differentially expressed in virulent isolates and low virulence isolates of L. major

mRNA was firstly purified from promastigotes from two highly virulent isolates (GLC94 and GLC67) and from two low virulence isolates (GLC32 and 07) then reverse transcribed to cDNA using Oligo$(dT)_{11}$MN primers, where M = A or C or G and N = A or C or G or T. The primer used during the differential display experiments was:

The amplification reactions were carried out by PCR using the same oligo-dT primers used during the reverse transcription reaction and combined with 10 arbitrary primers, as described in the scientific literature (Liang and Pardee, 1992; Liang, Bauer et al, 1995; Heard, Lewis et al, 1996).

In total, 60 combinations of primers were produced and analyzed. Polyacrylamide gel analysis of the amplification products using different combinations of primers showed that the genes from different isolates from L major (highly virulent or low virulence) express, in approximately 95% of cases, the same mRNA and at equivalent levels. Taken alone, 25 messengers appear to be differentially expressed between highly virulent and low virulence isolates (Figure 1A). The differentially expressed cDNA was firstly isolated from the acrylamide gel then reamplified using the same combinations of primer used during the first PCR and finally cloned in pMOS vector. Sequencing the different clones showed that a certain number of them were identified.

Analysis of the mRNA from the different isolates of L major by Northern Blot using the 14 fragments differentially expressed as a probe showed that 3 clones out of the 14 isolates exhibit differential expression between highly virulent isolates and low virulence isolates. One of these clones, p14, has been characterized by Northern Blot. The probe corresponding to clone p14 specifically hybridized with a transcript with an approximate size of 2.2 kb, which is preferentially expressed in the two most virulent isolates compared with the two least virulent isolates (Figure 1B). This confirms the results obtained by the Differential Display technique. Clone p14 has been completely sequenced and the size of this clone is 339 bp. A comparison of

the nucleotide sequence of this fragment with the sequences described in the databases (GenBank and EMBL) did not identify a significantly homologous sequence. This could be due to the fact that the p14 clone corresponds to the non translated 3' terminal region of the messenger.

2- Cloning and analysis of the entire cDNA p14 sequence

To isolate the entire cDNA sequence corresponding to the p14 clone, the 339bp fragment was used to screen a cDNA bank of promastigotes of the GLC94 isolate. Two positive clones were isolated from $6X10^5$ recombinant clones analyzed. Figure 2 shows the nucleotide sequence of the longest clone, which is 2094bp (SEQ ID No: 1). This clone has an open reading frame coding for a 477 amino acid (aa) polypeptide with a theoretical molecular weight of 52.4 kDa and an isoelectric point of 5.22. The N-terminal region of this protein corresponds to a potential peptide signal for export to the endoplasmic reticulum, 20 aa long. The non translated 5' region contains a splice leader sequence characteristic of Leishmanias and the non translated 3' region contains a poly A tail preceded by a potential polyadenylation site (Figure 2).

The peptide sequence for the isolated clone showed 27-36% identity with proteins of the protein disulfide isomerase family (PDI and Erp) of several species (Figure 3). Further, this protein contains two regions at residues 47—52 and 381-386 which are identical to the potential active sites (Cys-Gly-His-Cys, or CGHC) of PDI, Erp and proteins from the thioredoxin family. The C-terminal portion shows a potential signal for retention in the endoplasmic reticulum of the KDEL (EEDL) type at residues 474-477 suggesting that, like PDI and Erp, this protein is found in the cavity of the endoplasmic reticulum. P14 is thus a protein from the L major protein disulfide isomerase family. It has been denoted LmPDI (Figure 2 and 3).

To determine whether LmPDI is endowed with an oxido-reductase thiodisulfide activity as demonstrated for the majority of the protein disulfide-isomerase described, the capacity of the recombinant protein LmPDI to renature denatured RNase A was studied. Recombinant LmPDI protein was synthesized in E coli than purified and used in a test, in vitro, for reactivating RNase.

The results obtained show that LmPDI is capable of restoring RNase A activity in a similar manner to that of bovine PDI, used as a control (Figure 4).

To identify the number of copies of the gene coding for LmPDI, the inventors carried out Southern Blot type hybridization using as a probe the cDNA fragment of $^{32}P$ labeled LmPDI. The results obtained generally showed a single band, except for enzymes with a cleavage site within the cDNA of LmPDI (Figure 5A). The gene coding for LmPDI is thus probably present in a single copy in the genome for L major. Further, the gene for LmPDI appears to be conserved in different species of the Leishmania tested (Leishmania infantum, dermotropic, and a viscerotrope, Leishmania donovani) Figure 5B).

3-Immunoblot analysis of LmPDI expression

To characterize the expression of the native protein, a rabbit was immunized with recombinant LmPDI protein synthesized in E coli and purified by affinity chromatography. Using immunoblot, the inventors have shown that the anti-LmPDI polyclonal antibody obtained strongly recognized a protein of the expected size (55 kDa) in lysates from promastigotes in the stationary growth phase of GLC94 (Figure 6). Further, two other proteins were detected. The first had a molecular weight of 105 kDa, corresponding to about twice that of LmPDI, and the second had a molecular weight of 35 kDa. In order to verify whether the 105 kDa protein corresponded to a dimer of LmPDI, denatured GLC94 promastigote lysates were analyzed in the presence of high concentrations of DTT (0.5 mM). Under these conditions, the anti-LmPDI detected no more proteins of 105 kDa. These results suggest that LmPDI is organized into oligomers. The 35 kDa protein appears to be a contaminant. In fact, anti-LmPDI purified on an affinity column (Sepharose 4B-LmPDI) no longer recognizes the 35 kDa protein (Figure 6).

In order to compare the level of expression of LmPDI between the most and the least virulent isolates, promastigote proteins were extracted then quantified in the stationary growth phase. 5 µg of proteins were analyzed on a 12% polyacrylamide-SDS gel and transferred to a nitrocellulose membrane. Western blot analysis using an anti-LmPDI antibody showed that

LmPDI (55 kDa) and its dimer (105kDa) were more strongly expressed in the most virulent isolates (Figure 7). In contrast, the 35 kDa contaminating protein was expressed in an equivalent manner regardless of the test strain. These results suggest a correlation between the level of expression of LmPDI and the pathogenic power of the studied isolates.

**Example 3: Induction by LmPDI of in vitro proliferation of mononuclear cells from individuals having active lesions or ZCL antecedents**

L. major LmPDI, because of its high expression during the infectious stage of the parasite, could be a target for a cellular immune response. In order to verify the pertinence of this hypothesis, the capacity of LmPDI to induce a cellular immune response was evaluated by means of experiments on the proliferation of mononuclear cells obtained from individuals having active lesions or ZCL antecedents.

This study was carried out in 37 individuals living at El Guettar (southern Tunisia) for whom the results of the cellular proliferation test against total antigens from the parasite (SLA, a test indicating prior contact with the parasite) was available. These individuals were divided up as follows:

Group 1: composed of 8 individuals with a negative SLA test;

Group 2: composed of 29 individuals with a positive SLA test.

Mononuclear cells comprising lymphocytes and monocytes were separated from peripheral blood by centrifuging on a Ficoll/Hypaque gradient (Pharmacia, Uppsala, Sweden).

The results (Figure 8) show significant proliferation with immune individuals.

Cytokines (IFN-β, IL-4) in PBMC culture supernatants were induced by incubating mononuclear cells for 48 hours with the same concentration of LmPDI and assaying by means of an ELISA test using human anti-IL-4 and anti-IFN-β monoclonal antibodies (Pharmingen, San Diego, CA).

The results came from a small sample of individuals. They clearly show the absence of IL-4 and the presence of significant amounts of IFN-β in the supernatant from cells stimulated by LmPDI.

This result shows an essentially type Th1 response, indicating that LmPDI could constitute a vaccine candidate against Leishmania.

**Example 4: Inhibition of growth of Leishmania major in a liquid medium in the presence of a PDI inhibitor**

Bacitracin is a known PDI inhibitor. Experiments using bacitracin showed that at the final concentration of 2 mM, bacitracin completely inhibited the growth of L. major parasites in a liquid medium (Figure 9).

These experiments were carried out under the following experimental conditions:

a) Preparation of scrambled RNase A:

20 mg of purified ribonuclease (RNase A) was reduced and denatured at ambient temperature for 18 hours in a buffer containing 0.15 M of DTT, 6M of guanidine-HCl and 0.1 M of Tris-HCl at a pH of 8.6, before being purified on a Sephadex G25 column equilibrated in 0.01 M HCl. The concentration of scrambled RNase A fractions was determined with the help of an extinction coefficient of 9200 $M^{-1}cm^{-1}$ at 275 nm. The fractions were stored at -80°C for two weeks.

b) Reactivation of RNase A in the presence of recombinant LmPDI protein

The scrambled RNase A (8 µM) was incubated in a buffer containing 4.5 mM cCMP, 1 mM glutathione GSH, 0.2 mM glutathione disulfide GSSH, 2 mM EDTA and 100 mM Tris-HCl, pH 8, in the presence of bovine serum albumin (BSA) (1.4 µM) as a negative control, bovine protein disulfide-isomerase (1.4 µM) as a positive control, or recombinant LmPDI (1.4 µM) for 30 minutes at 25°C. RNase A reactivation was determined by measuring the RNase A activity at 296 nm every 5 minutes for 30 minutes as described in the literature (Lyles and Gilbert, 1991).

c) In vitro tests for inhibition of the thiodisulfide oxido-reductase activity of recombinant LmPDI by different PDI inhibitors

The experimental conditions for the thio-disulfide oxido-reductase activity inhibition tests for recombinant LmPDI were strictly identical to those described in the paragraph "Reactivation of RNase A in the presence of recombinant LmPDI protein", except that the reactions were carried out in the presence of 0.01 mM, 0.1 mM, 0.5 mM and 2 mM of the following PDI inhibitors:

- bacitracin;
- zinc bacitracin;
- p-chloromercuribenzoic acid (pCMBA);
- tocinoic acid.

d) Inhibition of parasite (L. major) growth in a liquid medium

With the aim of determining the effect of bacitracin (BAC), zinc bacitracin (BACZn), p-chloromercuribenzoic acid (pCMBA) and 5,5'-dithiobis(2-nitrobenzoic acid) (DTNB) on the in vitro growth of Leishmanias in a liquid medium, different concentrations of the inhibitors cited above, 0 mM, 0.05 mM, 0.1 mM, 0.2 mM, 0.5 mM, 1 mM, 1.5 mM, 2 mM, 2.5 mM and 5 mM were added to RPMI supplemented with 5% fetal calf serum and containing $2 \times 10^6$/ml of parasites in the exponential growth phase. The parasites were incubated at 26°C and counted every 24 hours over 96 hours. The parasites were counted on Mallassez cells.

**Example 5: Evaluation of LmPDI inhibiting molecules as regards their capacities to stop the growth of Leishmania major**

The evaluation of the role of LmPDI in Leishmania virulence allows novel strategies for identifying molecules that are active in treating Leishmania to be envisaged. It is probable that molecules known for their anti-PDI activities other than bacitracin are capable of inhibiting the growth of the parasite. An example of a protocol for evaluating molecules potentially effective against Leishmania is presented here.

An evaluation of molecules known for their anti-PDI activities or those which will be newly identified can be carried out in three steps. During the first step, the molecules are tested in vitro on recombinant LmPDI protein produced in Escherichia (E.) coli. Tests for inhibiting the growth of parasites in a liquid medium are then carried out and finally, the molecules are tested in the experimental murine model of leishmaniasis.

1- Evaluation of the inhibition of recombinant LmPDI. The detailed technique for analyzing the PDI activity of LmPDI was described in Example 2 and in the Materials and Methods section above. The same technique can be used to evaluate the capacity of certain PDI inhibitors which are known or still to be identified (using a molecular model of LmPDI), by adding different concentrations of potential inhibitors to the reaction volume. The results are expressed as the percentage inhibition compared with the buffer alone. Only molecules with a significant and dose-dependent LmPDI inhibiting activity are retained.

With the aim of determining whether the different PDI inhibitors described in the literature could inhibit the thio-disulfide oxido-reductase activity of LmPDI, the capacity of these inhibitors to block the enzymatic activity of LmPDI synthesized in E. coli then purified was studied in an in vitro test, at different concentrations. The inhibitors were:

- bacitracin;
- zinc bacitracin;
- p-chloromercuribenzoic acid (pCMBA);
- tocinoic acid.

The results obtained are shown in Figure 10 and show that the LmPDI activity is completely inhibited in the presence of 0.01 mM of pCMBA and 2 mM of bacitracin or zinc bacitracin. In contrast, tocinoic acid did not appear to have a very great effect on the activity of LmPDI at the concentrations employed (concentrations which completely inhibit the activity of human PDI).

2- Inhibition of the growth of parasites (L. major) in a liquid medium. The test molecule was dissolved in a solvent the suitability of which depended on its physico-chemical properties (solubility in aqueous solutions or organic solvents). In all cases, the solvent alone was used as a control. As an example, the experiments could be carried out on the GLC94 L. major isolate. The composition of the culture medium was given above (Example 2 and Materials and Methods). The cultures were incubated at 26°C and re-pricked out regularly to maintain the parasites in the stationary growth phase. For certain experiments, the amastigote-like stage of the parasite was used. In this case, the parasites (promastigotes) of the stationary growth phase were centrifuged, the medium was replaced with Schneider Drosophila medium adjusted to a pH of 5.0 and supplemented with fetal calf serum (FCS). The cultures were then incubated under 5% $CO_2$ at 35°C.

Inhibition of the growth of L. major promastigotes was carried out on parasites taken in the exponential growth phase, adjusted to the initial concentration of $10^6$ parasites/ml of complete medium and incubated in an amount of 100 µl/well in 96-well culture plates in the absence or presence of different concentrations of the test molecule. The parasites were incubated in 5% $CO_2$ at 26°C and counted every 24 h for 96h. Parasite counting was carried out on a Mallassez cell. Alternatively, a hemocytometer could be used. All of the measurements were carried out in triplicate. The inhibiting capacity of a molecule was determined as the inhibiting concentration which reduces cell division by 50% compared with the control (IC50).

The reduction in the viability of amastigotes was evaluated using a fluorimetric test employing Almar Blue as a viability/growth indicator.

The PDI inhibitors described in paragraph (1-) were tested with the aim of evaluating their capacities to inhibit the in vitro growth of parasites. To this end, different quantities of inhibitors were added to RPMI containing $2 \times 10^6$/ml of parasites in the exponential growth phase. The parasites were incubated at 26°C and counted every 24 hours over 96 hours. The parasites were counted on Mallassez cells. The results obtained are shown in Figure 11 and

show that bacitracin, zinc bacitracin or pCMBA completely inhibited the growth of leishmania in concentrations of 5 mM and 2 mM and 0.5 mM respectively. In contrast, 5,5'-dithiobis(2)nitrobenzoic acid) (DTNB) in the concentrations used (concentrations which completely inhibit the activity of human PDI) did not appear to have a very large effect on the growth of Leishmanias.

3- Evaluation of the efficacy of pre-selected inhibitors in the experimental model of infection of sensitive BALB/c mice by L. major. The in vivo experiment will depend on the toxicity and physico-chemical properties of the test molecules. BALB/c mice will be infected by $10^6$ L. major promastigotes obtained during the stationary growth phase and injected (in a volume of 50 µl) into the plantar pad of the rear right paw. The lesion diameter will be measured weekly using sliding calipers.

In all, three therapeutic protocols will be applied depending on the case:

- for hydrophobic molecules, which diffuse well, and are slightly toxic or non-toxic, the product will be injected intraperitoneally at different concentrations and using different schemes. The frequency of injection will depend on the bioavailability of the molecule and on its half-life. In all cases, the protocol will be stopped at the end of 9 weeks following infection;
- for hydrosoluble and relatively toxic molecules, the injections will be made intra-lesionally (in general, the active doses can be divided by 10) by dint of at least four injections into the indurated zone;
- for liposoluble molecules, a pomade will be tested by weekly application to the experimental lesion.

Overall, and regardless of the mode of injecting the test product, two types of protocols will be carried out:

- a protocol which starts immediately after injecting the parasites;

- a protocol which starts 4 to 5 weeks after injecting parasites, at a time at which the lesion will already have been established.

In all cases, at the end of the protocol, the mice will be sacrificed and an estimate of the parasitic load will be made at the injection site and in the ganglion which drains the lesion.

**Example 6: In vitro infection for murine macrophages by Leishmania**

Murine bone marrow macrophages (MBMM) were obtained from bone marrow extruded from a femur or tibia from female BALB/c mice. The MBMM was cultivated in multi-chamber plates in an amount of $1.5 \times 10^3$ cells per well in 500 µl of complete medium. To stimulate the growth and maturation of the MBMM, the culture medium was supplemented with 20% of medium conditioned with L-929 fibroblasts as a source of macrophage colony stimulating factor (MCSF). After 6 days of culture at 37°C and 5% $CO_2$, the medium was removed, the MBMM was washed, and fresh RPMI medium with 10% fetal calf serum but comprising no medium conditioned by L-929 fibroblasts was added. Intra-lesional amastigotes were purified from non-ulcerated lesions by differential centrifugation and counted using trypan blue viral stain. These parasites were used to infect the MBMM in a final ratio of four parasites per macrophage. Two hours after adding the parasites, the macrophages were washed five times with PBS to eliminate non phagocytary amastigotes. The cultures were then incubated at 37°C in 95% air and 5% $CO_2$. The experiments were carried out at different points in time: 30 minutes and 2, 24 and 72 hours. At the indicated times, the wells were rinsed with PBS, the covers were removed and the infected macrophages were fixed with ethanol for 1 hour at ambient temperature. The plates were then washed and stained with Giemsa to follow the infection.

The infected macrophages were counted in the centre of each well were the cells were well spread out and the parasites could be counted easily. At this level of the plate, the parasite/macrophage ratio could be more than 4.

**Example 7: Inhibition of the enzymatic activity of recombinant LmPDI by protein disulfide-isomerase inhibitors**

Several protein disulfide-isomerase (PDI) inhibitors have been described in the literature (Ryser et al, 1994, Orlandi 1997, Mou et al, 1998). Of these, bacitracin and zinc bacitracin constitute a complex of polypeptide antibiotics produced by Bacillus subtilis and Bacillus lichenformis. Bacitracin A is the principal compound of commercial bacitracin, which is a mixture of at least nine bacitracins. This antibiotic is capable of inhibiting synthesis of the wall of many Gram+ bacteria, but also the activity of many proteases such as PDI, transglutaminase, papain and neuropeptidase. The majority of those proteases have a cysteine residue in their active site.

In a first step, the inventors tested the effect of these inhibitors in verifying their possible ability to alter the enzymatic activity of recombinant LmPDI (rLmPDI) in vitro.

The scrambled RNase technique described above (Lyles and Gilbert, 1991) was used to demonstrate the activity of LmPDI. Twenty milligrams of RNase A (Amersham-Pharmacia) was denatured in a buffer composed of 0.15 M dithiothreitol, 6M guanidine HCl and 0.1 M Tris-HCl, pH 8.6 for 18 hours at ambient temperature. The scrambled RNase was then purified on a Sephadex G25 column equilibrated in HCl 0.01 M and quantified by spectrophotometry at 275 nm.

In a glutathione-based reducing buffer, PDI catalyzes renaturing of scrambled RNase (Gilbert, 1998). Restoration of RNase activity was measured by spectrophotometry in the presence of cytidine 2'-3'-cyclic monophosphate (cCMP) as a substrate. 8 µM of scrambled RNase, alone or in the presence of 1.4 µM of bovine serum albumin (BSA) or 1.4 µM of rLmPDI was mixed in a buffer containing 4.5 mM of cCMP, 1 mM of reduced glutathione (GSH), 200 µM of oxidized glutathione (GSSG), 2 mM EDTA and 100 mM Tris-Cl, pH 8. The reaction was carried out at ambient temperature for 30 minutes. The hydrolysis of cCMP resulting from the renaturing of RNase was recorded by measuring the absorbance at 296 nm every 5 minutes for the half hour of the reaction.

The activity of the recombinant LmPDI (rLmPDI) was measured in the presence of different concentrations of bacitracin (BAC 0.01 mM – 2 mM) and zinc bacitracin (BACZn, 0.01 mM – 2 mM). The results are shown in Figure 12.

These results show that bacitracin and zinc bacitracin have similar effects. In the presence of these two products, 50% inhibition was observed at 0.1 mM, 70% at 0.5 mM and 100% at 2 mM. The concentrations which inhibit rLmPDI are comparable with those described in the literature as inhibitors for PDIs from other species.

**Example 8: In vitro growth kinetics of L. major promastigotes in the presence of zinc bacitracin**

The inventors then tested the effect of zinc bacitracin on the in vitro growth kinetics of L. major promastigotes. For this study, only zinc bacitracin was tested, firstly because it had the same rLmPDI enzymatic activity inhibition profile as bacitracin, and secondly because bacitracin is more stable and less toxic when coupled with zinc.

To this end, promastigotes from the GLC94 isolate were cultured on a medium based on coagulated rabbit serum for two days. Then the parasites ($2 \times 10^6$ parasites per ml) were transferred into complete medium comprising zinc bacitracin BACZn, in concentrations of 1, 1.5 and 2.5 mM. Promastigotes cultured in complete medium in the absence of inhibitors were used as the control. Monitoring was by counting the parasites at 48, 72 and 96 hours. The results are shown in Figure 13.

These results show that zinc bacitracin partially inhibits the growth of parasites at 1.5 mM with complete inhibition at 2 mM and at 5 mM, while it had no effect at 1 mM. Thus, it is very important to note that this molecule is capable of blocking the proliferation of L major parasites in culture.

**Example 9: Inhibition of the growth of L. major promastigotes in BALB/c mice in the presence of zinc bacitracin**

The availability of zinc bacitracin, which already forms a weapon in the therapeutic arsenal, has allowed it to be tested on the evolution of infection in the BALB/c mouse with L. major. Mice were infected with promastigotes in the stationary growth phase ($10^6$ promastigotes per paw) of the GLC94 isolate into the plantar pad of the rear paw and treated with a pomade based on 5 mM or 25 mM of BACZn (prepared in Vaseline). Treatment with the pomade was started 48 hours after injecting the parasites, by dint of one application per day over 5 days of the week. Mice infected in the same manner and treated with Vaseline were used as the control. The lesion size was measured each week. The results are shown in Figure 14.

Although preliminary, these results show that zinc bacitracin attenuates the progress of the disease when it is applied locally in the form of a pomade, at the injection site, to BALB/c mice. It should be emphasized that in the group of treated mice, lesion attenuation was observed in 2 out of 3 mice treated with 5 mM bacitracin and 2 out of 4 mice treated with 25 mM bacitracin. Recurrence of the clinical disease after stopping the treatment was expected since BALB/c mice are incapable of completely eliminating the parasite and even the treatments used in man (glucantime and paramomycin) have little effect on the disease induced in the BALB/c mouse, in which complete disappearance of the parasites has never been observed.

LmPDI can thus be considered to be a potential target for anti-leishmania chemotherapy and it appears that bacitracin is potentially effective against L major.

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